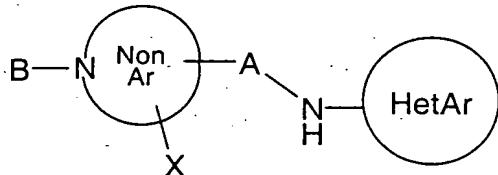


WHAT IS CLAIMED IS:

1. A compound having the formula (I):

5



(I)

or pharmaceutically acceptable salts thereof, wherein

10 NonAr is a nonaromatic 5-7 membered ring containing 1 or 2 nitrogen ring atoms or an aza bicyclo octane ring;

HetAr is a 5 or 6 membered heteroaromatic ring containing 1-3 nitrogen ring atoms, or isoxazolyl, thiazolyl, thiadiazolyl, quinolinyl, quinazolinyl, purinyl, pteridinyl, benzimidazolyl, pyrrolopyrimidinyl, or imidazopyridinyl;

15 HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-;

20 A is -C₀₋₄alkyl-;

B is aryl(CH₂)₀₋₃-O-C(O)-, heteroaryl(CH₂)₁₋₃-O-C(O)-, indanyl(CH₂)₀₋₃-O-C(O)-, aryl(CH₂)₁₋₃-C(O)-, aryl-cyclopropyl-C(O)-, heteroaryl-cyclopropyl-C(O)-, heteroaryl(CH₂)₁₋₃-C(O)-, aryl(CH₂)₁₋₃-, heteroaryl(CH₂)₁₋₃-, aryl(CH₂)₁₋₃-NH-C(O)-, aryl(CH₂)₁₋₃-NH-C(NCN)-, aryl(CH₂)₁₋₃-SO₂-,

25 heteroaryl(CH₂)₁₋₃-SO₂-, wherein any of the aryl or heteroaryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro; and

X is H, OH, F, C₁₋₄alkyl, C₁₋₄alkoxy, NH₂, or X taken with an adjacent bond is =O.

2. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

5 NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

Sub A1
B is aryl(CH₂)₀₋₃-O-C(O)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

10 3. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom;

15 HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

20 4. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is an isoxazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

30 5. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is a thiadiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

~~4alkyl)(C0-4alkyl), nitro, (C1-2alkyl)(C1-2alkyl)NCH₂-, (C1-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.~~

~~5 6. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein~~

~~HetAr is a 5 membered heteroaromatic ring containing 2 nitrogen ring atoms;~~

~~HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C1-4alkyl, C1-4alkoxy, C2-4alkynyl, trifluoromethyl,~~

~~10 hydroxy, hydroxyC1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C0-4alkyl)(C0-4alkyl), nitro, (C1-2alkyl)(C1-2alkyl)NCH₂-, (C1-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.~~

~~15 7. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein~~

~~HetAr is quinolinyl optionally substituted with 1 or 2 substituents, each substituent independently is C1-4alkyl, C1-4alkoxy, C2-4alkynyl, trifluoromethyl, hydroxy, hydroxyC1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl,~~

~~20 cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C0-4alkyl)(C0-4alkyl), nitro, (C1-2alkyl)(C1-2alkyl)NCH₂-, (C1-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.~~

~~25 8. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein~~

~~HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C1-4alkyl, C1-4alkoxy, C2-4alkynyl, trifluoromethyl, hydroxy, hydroxyC1-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl,~~

~~30 cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C0-4alkyl)(C0-4alkyl), nitro, (C1-2alkyl)(C1-2alkyl)NCH₂-, (C1-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.~~

~~9. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein~~

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

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10. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is thiazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

11. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

12. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyrrolopyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

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13. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

5 HetAr is a imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl), nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

10

14. The compound according to Claim 2, or pharmaceutically acceptable salts thereof, wherein

15

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl), nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

20

15. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

25

B is aryl(CH₂)₁₋₃-SO₂-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁-4alkyl, C₃-6cycloalkyl, C₁-4alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

30

16. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

35

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl),

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nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

17. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is quinazolinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl), nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

18. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl), nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

19. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is imidazopyridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁-4alkyl, C₁-4alkoxy, C₂-4alkynyl, trifluoromethyl, hydroxy, hydroxyC₁-4alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀-4alkyl)(C₀-4alkyl), nitro, (C₁-2alkyl)(C₁-2alkyl)NCH₂-, (C₁-2alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

20. The compound according to Claim 15, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and

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HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), 5 nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

21. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

10 NonAr is a nonaromatic 5 membered ring containing 1 nitrogen ring atom; and

B is aryl(CH₂)₀₋₃-O-C(O)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

15

22. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atoms;

20

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

25

23. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

30 HetAr is pteridinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

35

24. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

10 25. The compound according to Claim 21, or pharmaceutically acceptable salts thereof, wherein

HetAr is benzimidazolyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

20 26. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

B is aryl(CH₂)₀₋₃-O-C(O)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

25

27. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 1 nitrogen ring atom; and

30 HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

28. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is purinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

10

29. The compound according to Claim 26, or pharmaceutically acceptable salts thereof, wherein

HetAr is a 6 membered heteroaromatic ring containing 2 nitrogen ring atom; and

15

HetAr is optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or

20 NH₂C(O)-.

30. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is an aza bicyclo octane ring; and

25

B is aryl(CH₂)₁₋₃-SO₂-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

30

31. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

35

B is heteroaryl(CH₂)₁₋₃-C(O)-, wherein the heteroaryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

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32. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

B is aryl(CH₂)₁₋₃—C(O)—, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

10 33. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

15 B is aryl—cyclopropyl—C(O)—, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

34. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

20 HetAr is pyridyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl—, phenylethynyl—, heteroarylethynyl—, —N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂—, (C₁₋₂alkyl)HNCH₂—, Si(CH₃)₃—C—, or

25 NH₂C(O)—.

35. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

30 HetAr is pyrazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl—, phenylethynyl—, heteroarylethynyl—, —N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂—, (C₁₋₂alkyl)HNCH₂—, Si(CH₃)₃—C—, or NH₂C(O)—.

36. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyridazinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

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10 37. The compound according to Claim 33, or pharmaceutically acceptable salts thereof, wherein

HetAr is pyrimidinyl optionally substituted with 1 or 2 substituents, each substituent independently is C₁₋₄alkyl, C₁₋₄alkoxy, C₂₋₄alkynyl, trifluoromethyl, hydroxy, hydroxyC₁₋₄alkyl, fluoro, chloro, bromo, iodo, cyano, methylsulfanyl, cyclopropylethynyl-, phenylethynyl-, heteroarylethynyl-, -N(C₀₋₄alkyl)(C₀₋₄alkyl), nitro, (C₁₋₂alkyl)(C₁₋₂alkyl)NCH₂-, (C₁₋₂alkyl)HNCH₂-, Si(CH₃)₃-C-, or NH₂C(O)-.

20 38. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

25 B is heteroaryl(CH₂)₁₋₃-O-C(O)-, wherein the heteroaryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro;.

30 39. The compound according to Claim 1, or pharmaceutically acceptable salts thereof, wherein

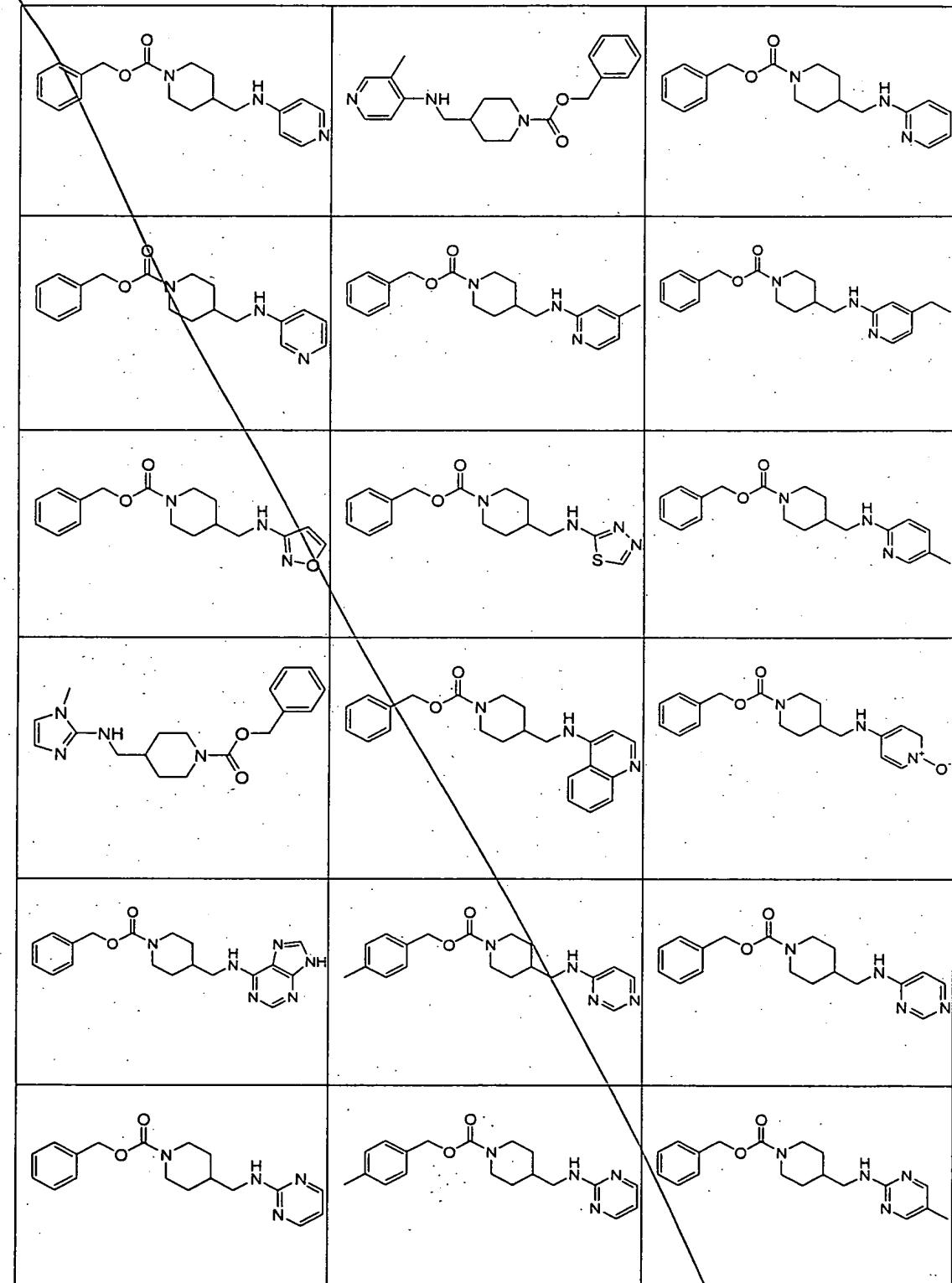
NonAr is a nonaromatic 6 membered ring containing 1 nitrogen ring atom; and

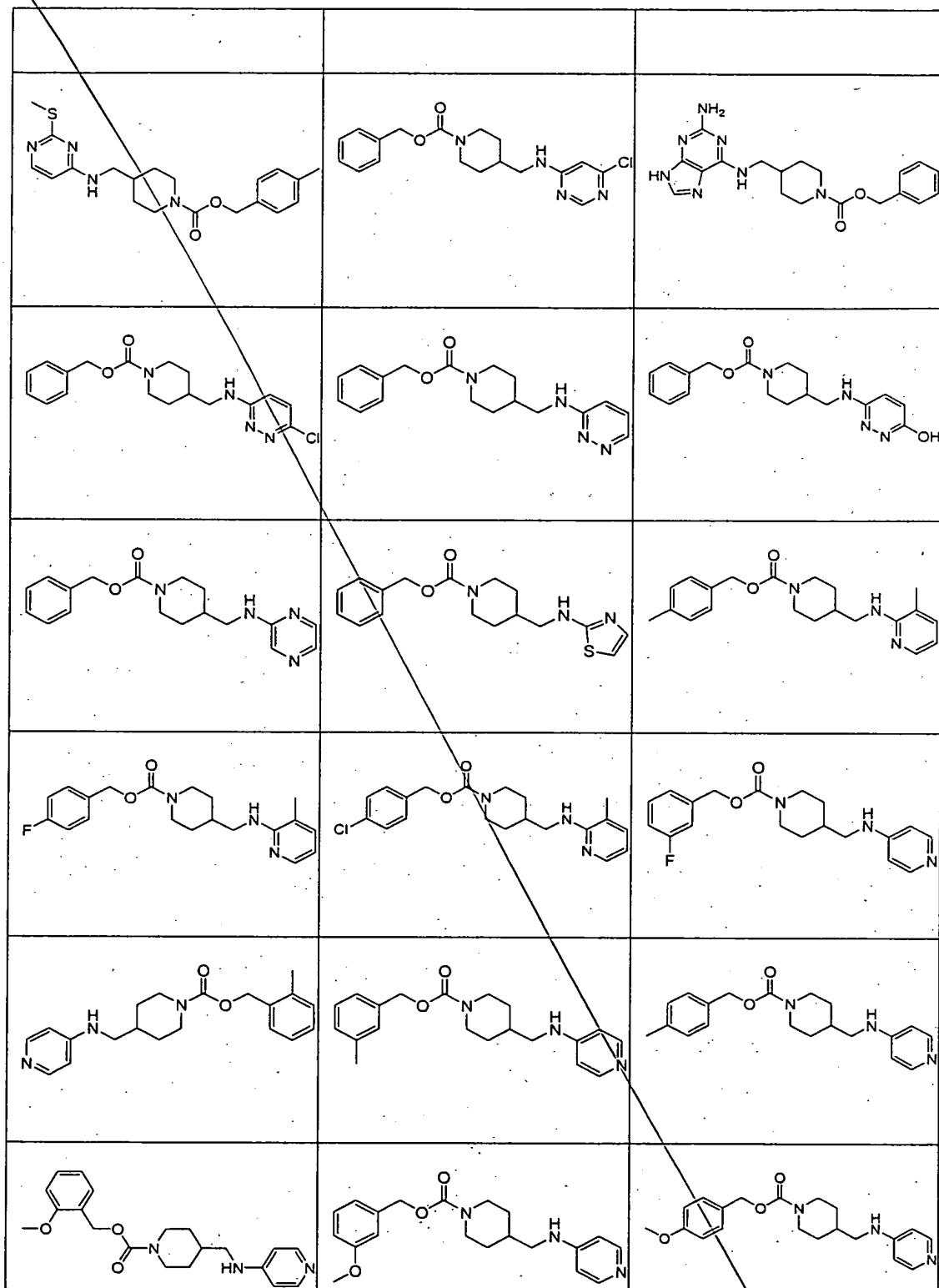
35 B is aryl(CH₂)₁₋₃-NH-C(NCN)-, wherein the aryl is optionally substituted by 1-5 substituents, each substituent independently is C₁₋₄alkyl, C₃₋₆cycloalkyl, C₁₋₄alkoxy, trifluoromethyl, bromo, fluoro, or chloro.

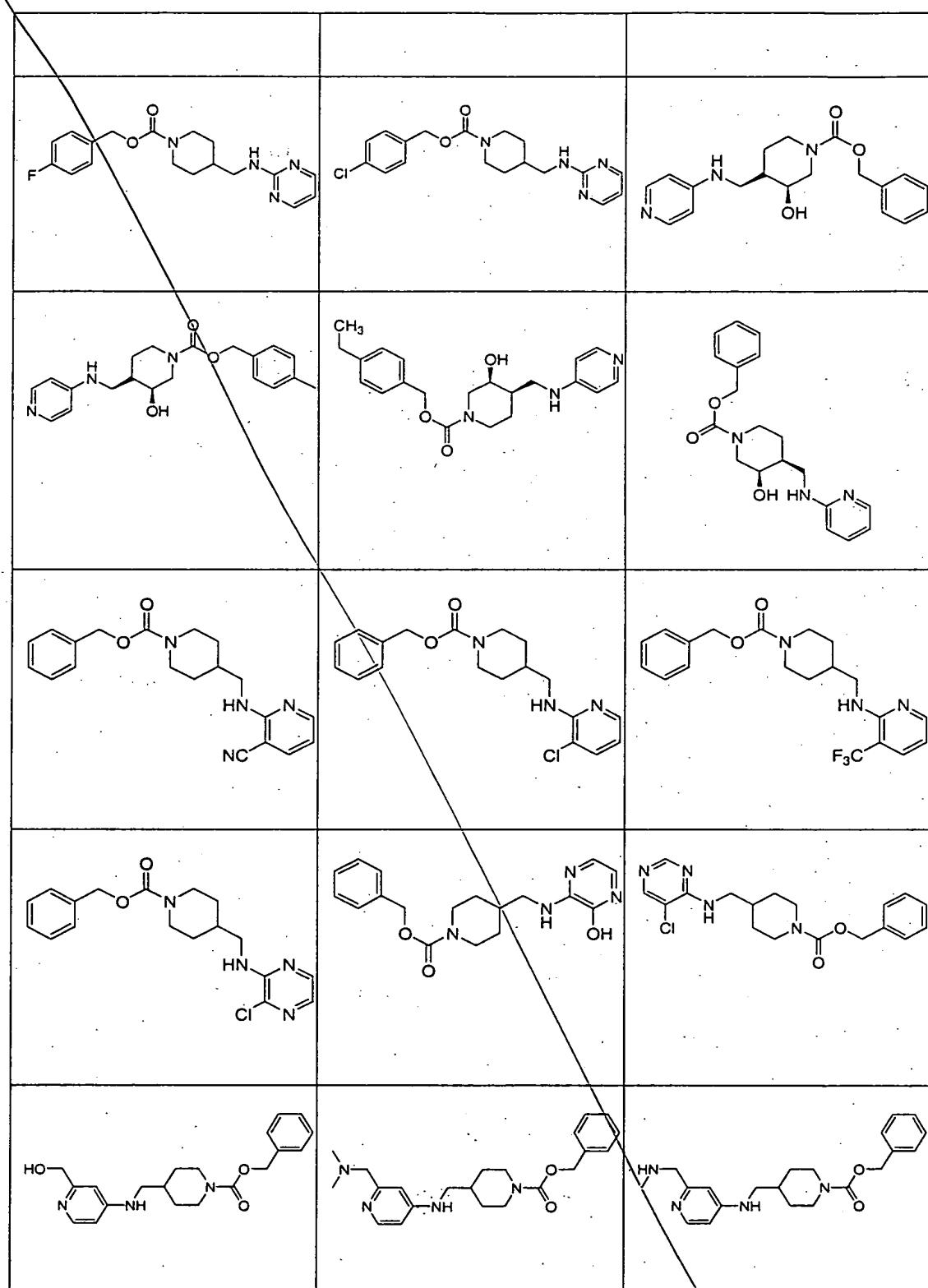
40. The compound according to Claim 1, wherein said compound is

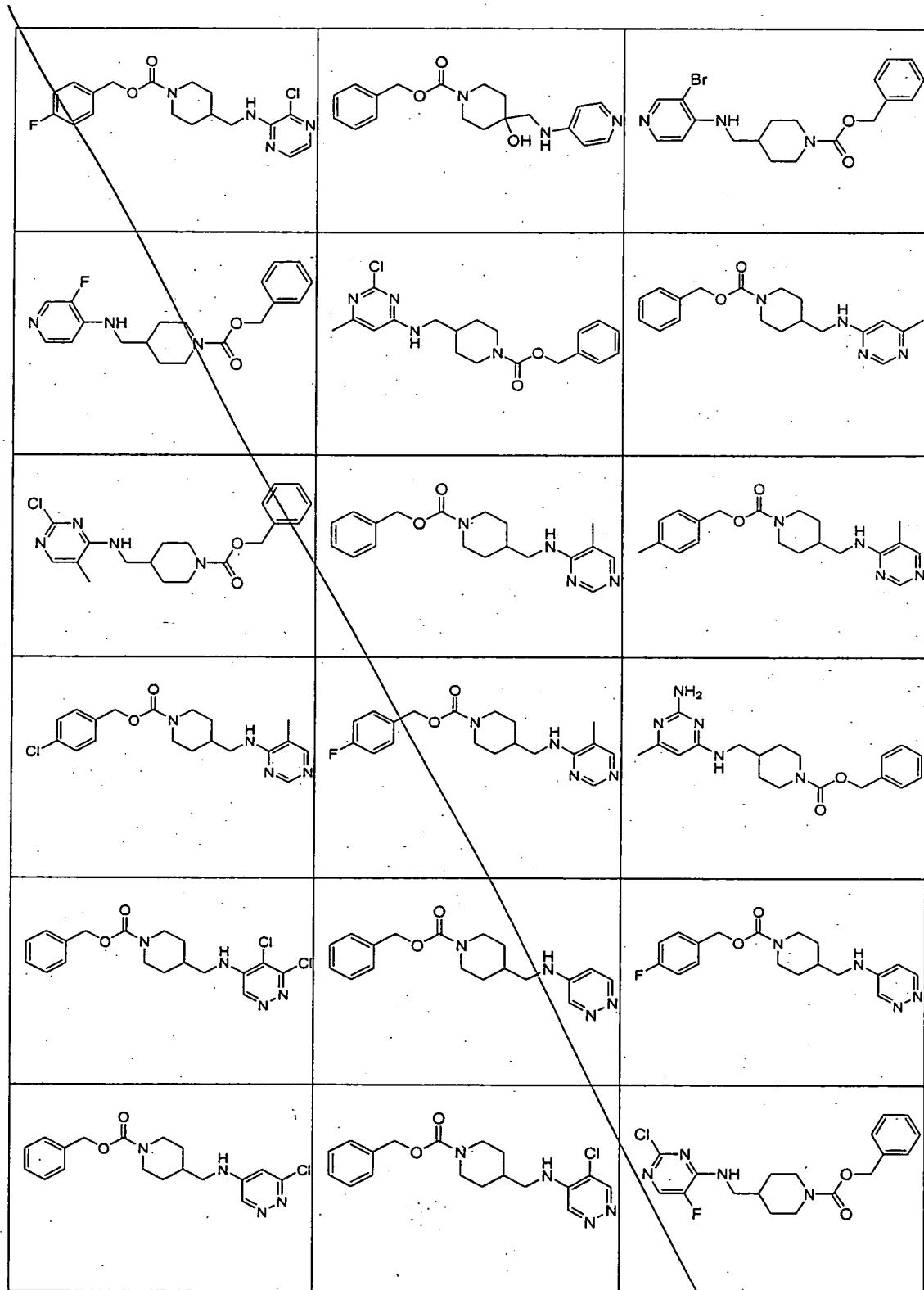
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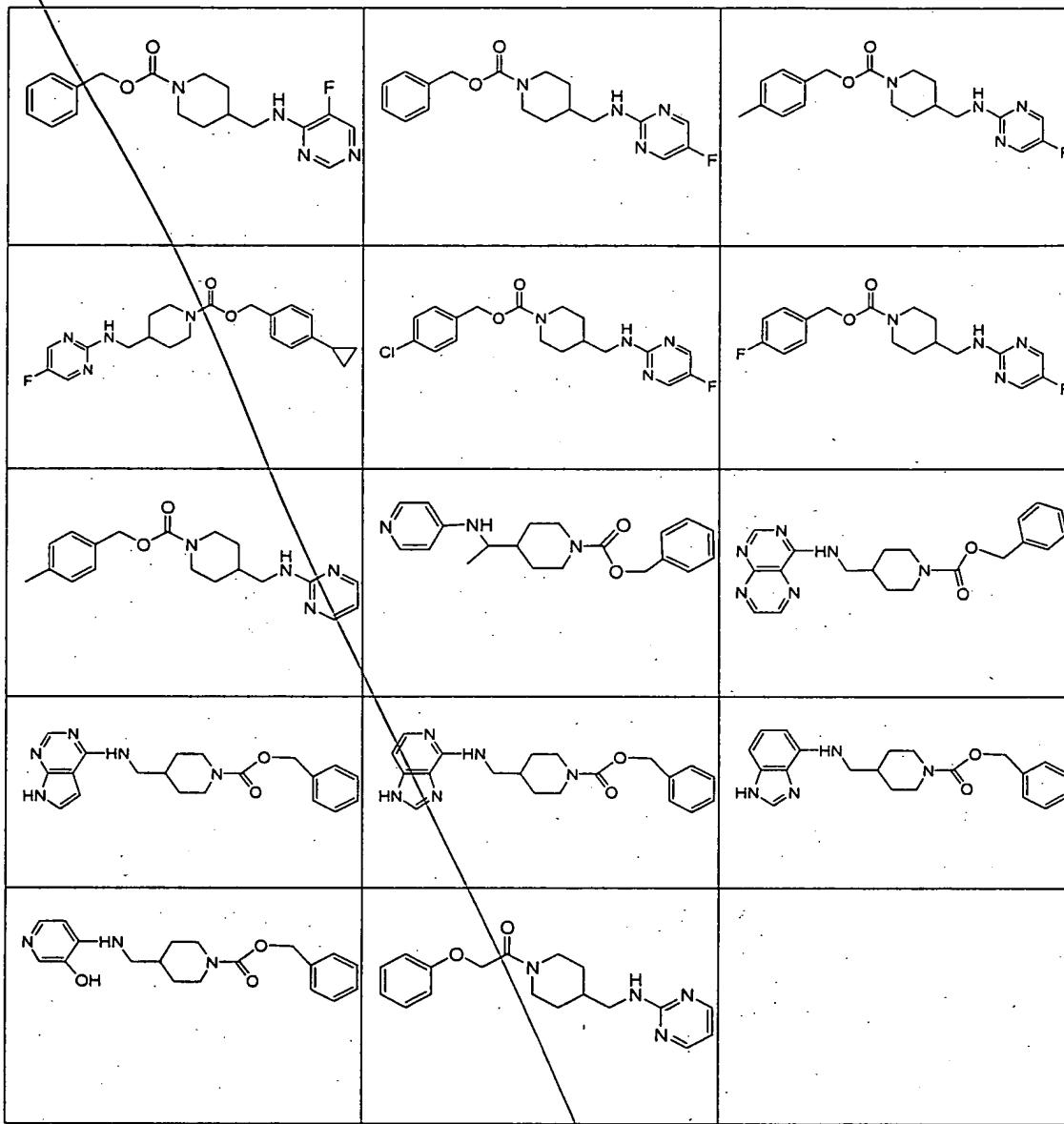
20832-200 20832-201 20832-202 20832-203 20832-204 20832-205



Sub
A1

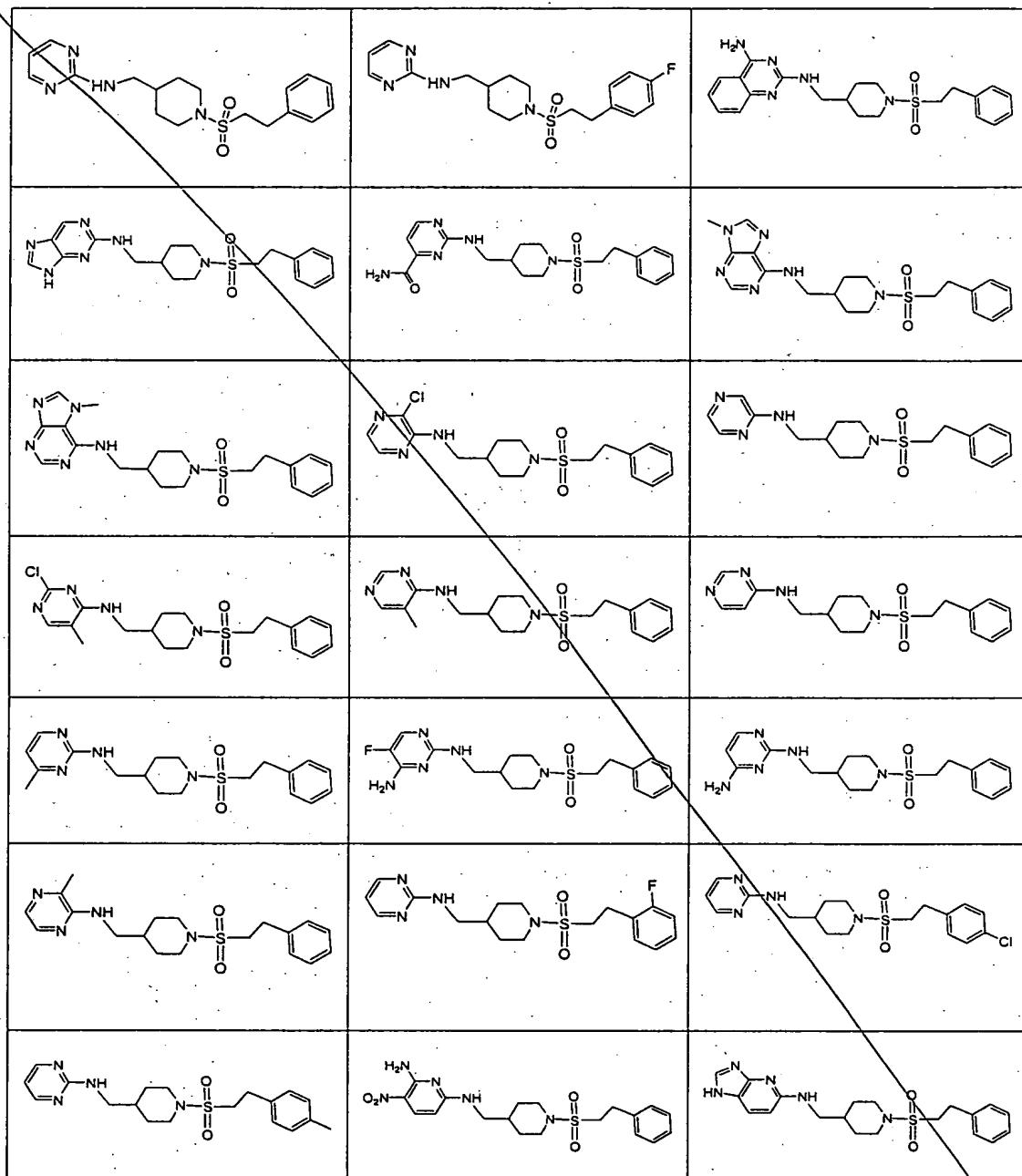
Sub
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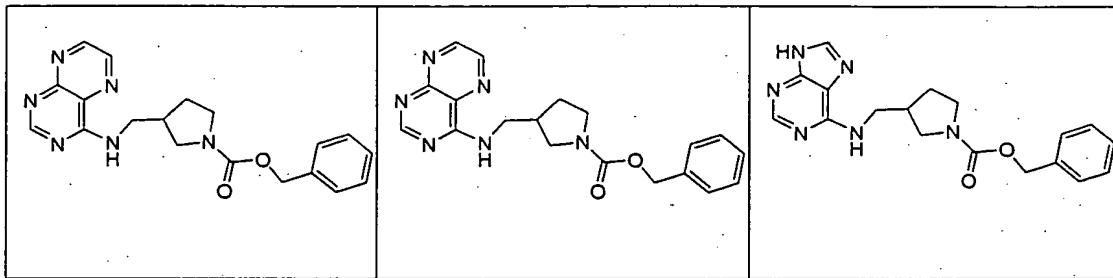
or a pharmaceutically acceptable salt thereof.

41. The compound according to Claim 1, wherein said compound is

Sub
A'

or a pharmaceutically acceptable salt thereof.

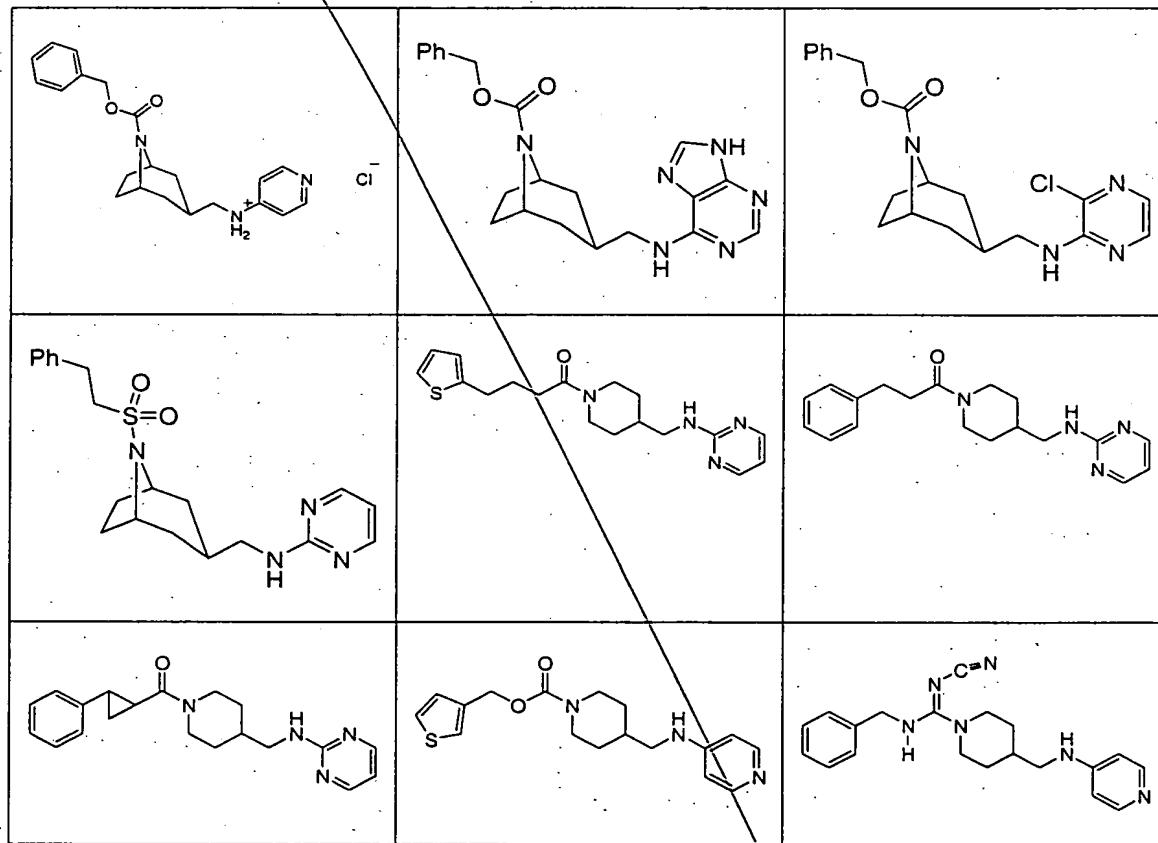
42. The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

43. The compound according to Claim 1, wherein said compound is

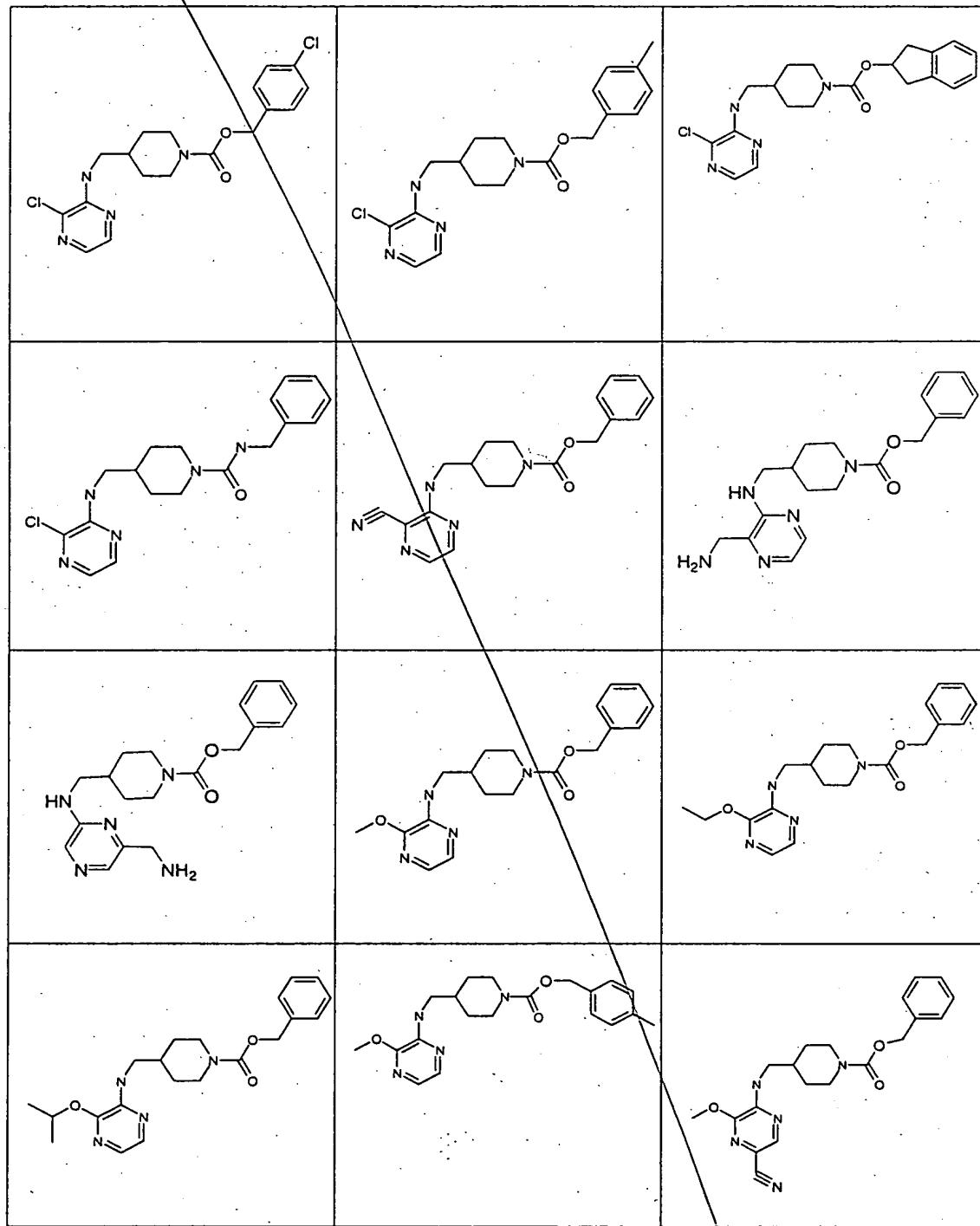
5

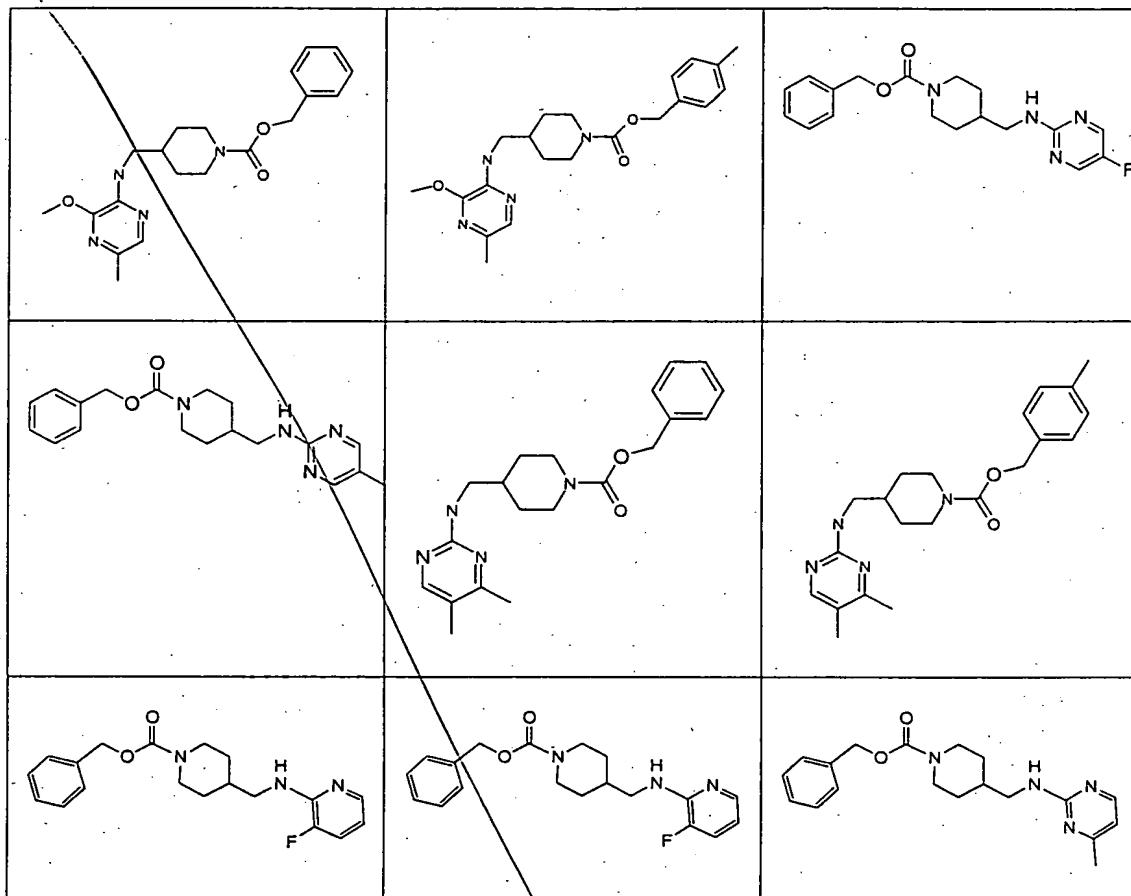


or a pharmaceutically acceptable salt thereof.

44. The compound according to Claim 1, wherein said compound is

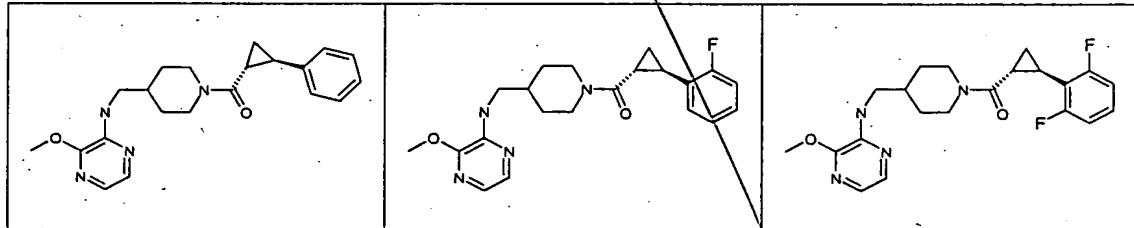
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A1



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A'

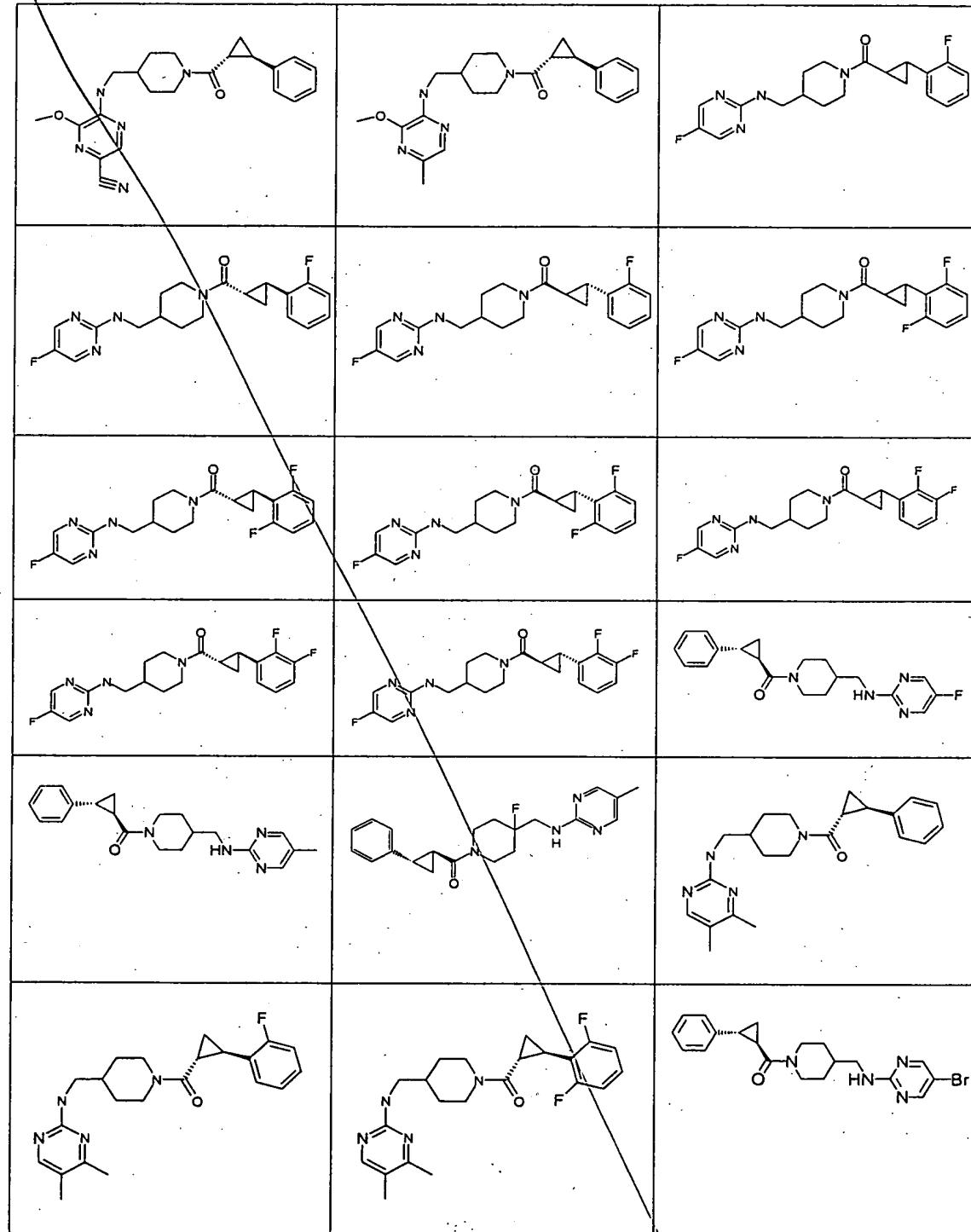
or a pharmaceutically acceptable salt thereof.

45. The compound according to Claim 1, wherein said compound is



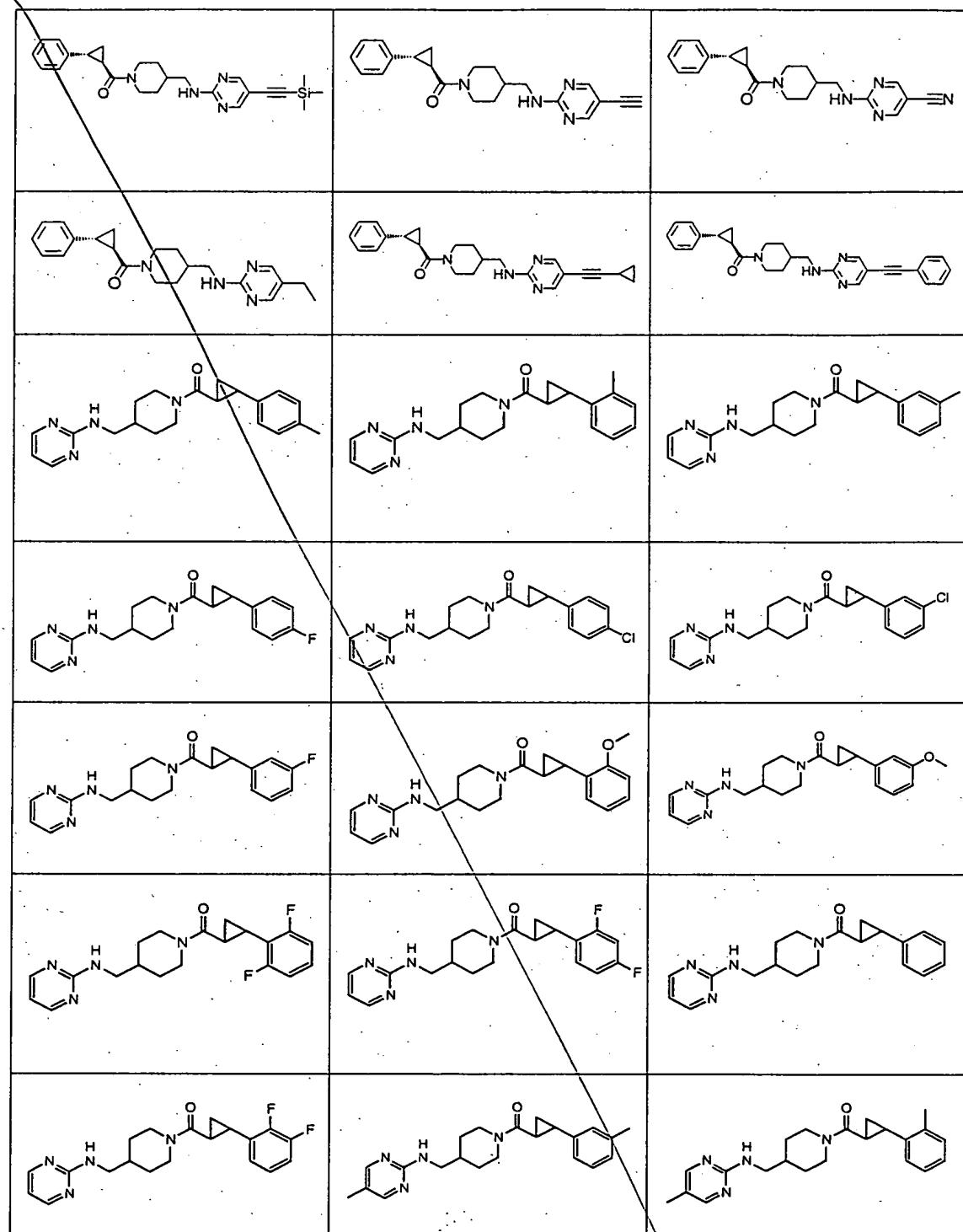
every day from 10 a.m. to 4 p.m. and from 6 p.m. to 8 p.m.

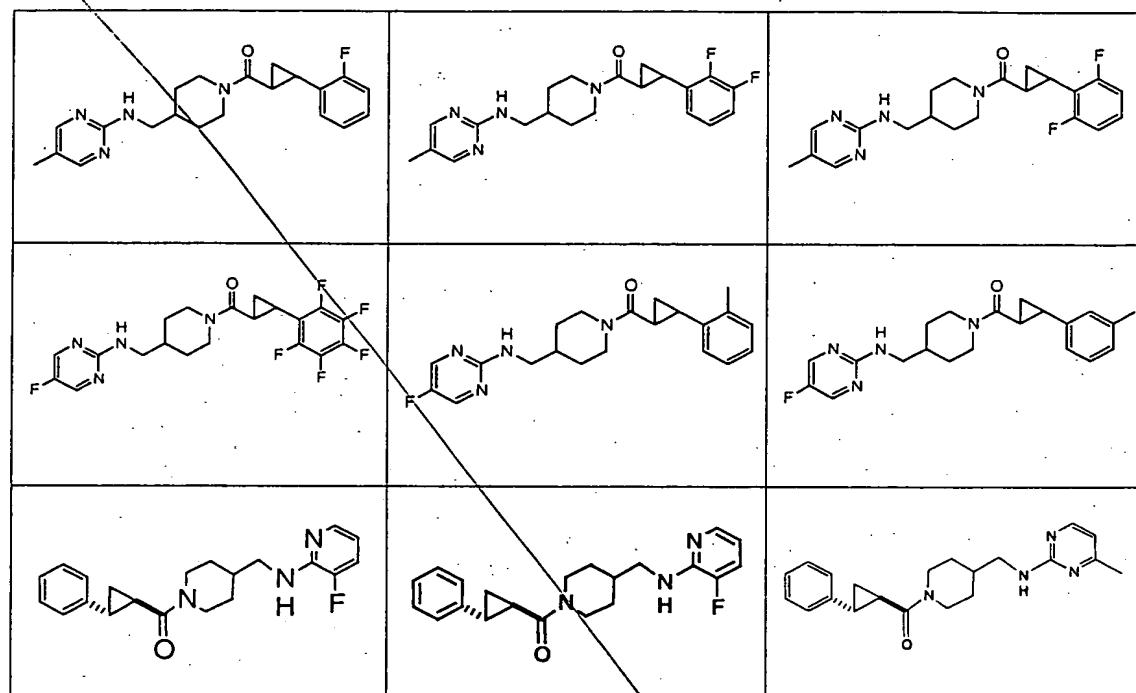
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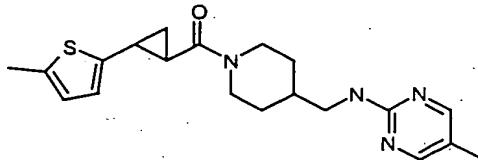
Such
all





or a pharmaceutically acceptable salt thereof.

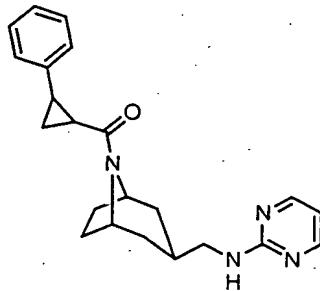
46. The compound according to Claim 1, wherein said compound is



5

or a pharmaceutically acceptable salt thereof.

47. The compound according to Claim 1, wherein said compound is



or a pharmaceutically acceptable salt thereof.

48. A pharmaceutical composition comprising an inert carrier and an effective amount of a compound according to claim 1.

49. The pharmaceutical composition according to claim 48 useful for the treatment of pain.

50. The pharmaceutical composition according to claim 48 useful for the treatment of migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke.

51. A method of treating pain comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.

52. A method of treating migraine, depression, anxiety, schizophrenia, Parkinson's disease, or stroke comprising a step of administering to one in need of such treatment an effective amount of a compound according to claim 1.